## **CLAIM AMENDMENTS**

1. (currently amended): <u>Process A process</u> for the preparation of an enantiomerically enriched compound of formula 1

$$R^{1}$$
 $R^{1}$ 
 $C^{*}$ 
 $R^{3}$ 
 $R^{4}$ 
 $R^{5}$ 
(1)

or a salt thereof, wherein:

C\* represents an asymmetric carbon atom;

R<sup>1</sup> and R<sup>2</sup> are different from each other, and, each independently, represent represents H, a substituted or unsubstituted alkyl or aryl group;

R<sup>3</sup> represents CH<sub>2</sub>OH or an optionally protected CHO group;

R<sup>5</sup> represents H, a substituted or unsubstituted alkyl or aryl group; and

 $R^4$  represents H or C(=O) $R^6$  wherein  $R^6$  represents H, a substituted or unsubstituted alkyl, aryl or alkoxy group or  $R^4$  represents or an amine protecting group, or  $R^4$  and  $R^5$  form together with the N to which they are attached a cyclic imide group, wherein

which process comprises hydrogenating an enantiomerically enriched compound with formula 2

$$R^{1} \xrightarrow{\stackrel{R^{2}}{\longrightarrow}} C = N \qquad (2)$$

$$NR^{7}R^{5}$$

or a salt thereof, wherein C\*, R<sup>1</sup>, R<sup>2</sup>, R<sup>3</sup> and R<sup>5</sup> are as defined above; and

 $R^7$  represents H or C(=O) $R^6$  wherein  $R^6$  is as defined above, or  $R^7$  represents an amine protecting group, or  $R^5$  and  $R^7$  form together with the N to which they are attached a cyclic [[imine]] imide group, is subjected to hydrogenation

in the presence of hydrogen, a hydrogenation catalyst and a mineral acid.

- 2. (currently amended): Process according to The process of claim 1, wherein R<sup>3</sup> is an optionally protected CHO group and wherein hydrogen is present at a hydrogen-pressure between 0.1 and 2 MPa.
- 3. (currently amended): Process according to The process of claim 2, wherein the hydrogen-pressure is between 0.5 and 1 MPa.

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- 4. (currently amended): Process according to anyone of claims 1-3 The process of claim 1 wherein R<sup>3</sup> is an optionally protected aldehyde and the amino aldehyde compound of formula 2 is isolated in the form of a chemically and configurationally stable derivative.
- 5. (currently amended): Process according to The process of claim 1, wherein R<sup>3</sup> is a CH<sub>2</sub>OH group and wherein at least during part of the hydrogenation hydrogen is present at a hydrogen-pressure between 2 and 10 MPa.
- 6. (currently amended): Process according to The process of claim 5, wherein at least during part of the hydrogenation the hydrogen-pressure is between 4 and 6 MPa.
- 7. (currently amended): Process according to The process of claim 5 [[or-6]], wherein the hydrogen-pressure initially is between [[0,5]] 0.5 and 2 MPa and subsequently, after most of the nitrile starting material compound of formula 2 is converted to the compound of formula 1, the hydrogen pressure is increased to a value between 2 and 10 MPa.
- 8. (currently amended): Process according to anyone of claims 1-7 The process of claim 1 wherein a Pd catalyst is used as the hydrogenation catalyst is a Pd catalyst.
- 9. (currently amended): Process according to anyone of claims 1-8 The process of claim 1, wherein as starting material, an enantiomerically enriched nitrile according to the compound of formula 2 is used that is prepared by

(precursor) fermentation, enzymatic resolution, crystallization induced asymmetric transformation, classical resolution, resolution via preferential crystallization, diastereomeric synthesis, catalytic asymmetric synthesis or dehydratation dehydration of at least one amino acid [[amides]] amide.

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